

## **Recent Actions to Lower Out-of-Pocket Costs on Drugs**

## **Congressional Actions**

Closing the Medicare Part D donut hole sooner and providing new discounts for innovative biosimilars. (H.R. 1892, Bipartisan Budget Act)

Lowering premiums for some by as much as 20 percent and repealing Obamacare provisions that drive up the overall cost of drugs. (<u>H.R. 1628</u>, the American Health Care Act)

Increasing generic competition to drive down the cost of drugs. (<u>H.R. 2430</u>, the <u>FDA</u> Reauthorization Act)

Modernizing the process for over-the-counter drugs to ensure patients have access to affordable therapies. (H.R. 5333, the Over-the-Counter Monograph Safety, Innovation and Reform Act)

Allowing more Americans to use pre-tax dollars for their healthcare expenses, including over-the-counter drugs. (<u>H.R. 6199</u>, the Restoring Access to Medication and Modernizing Health Savings Accounts Act)

## **Administrative Actions**

The Department of Health and Human Services released a <u>final regulation</u> to reduce drug costs for seniors enrolled in a Medicare Part D plan. The regulation allows for expedited generic substitution, applies generic cost-sharing to biosimilar and interchangeable biological products for low income Part D enrollees, and eliminates a provision that allowed plans to exclude a dedicated generic tier for the tiering exceptions process. All three of these policies help lower out-of-pocket costs for seniors.

The Administration announced its <u>Patients First</u> blueprint for lowering prescription drug prices. The plan calls for increasing competition, improving negotiation, incentives for lower list prices, and lowering out-of-pocket costs of prescription drugs.

The Administration proposed in the <u>physician fee schedule</u> to cut payments for new drugs from wholesale acquisition cost (WAC) + 6 percent down to WAC + 3 percent, decreasing the 20 percent out of pocket cost-sharing that patients are responsible for.

For Internal Purposes Only-Prepared by Ways and Means Committee Majority Staff

The Administration released its <u>Biosimilar Action Plan</u>. This plan lays out how the Administration intends to accelerate innovation in the biosimilars market.

Recently, the court ruled in favor of the Administration on a 340B <u>policy change</u> in the 2018 outpatient prospective payment system regulation, which is expected to will help Medicare beneficiaries save on coinsurance on Part B drugs administered at hospitals that participate in the 340B program by reducing the amount Medicare pays for those drugs. When utilized correctly, the 340B program allows hospitals caring for under-served and low-income populations to buy drugs at a lower cost in exchange. As a result of the Administration's 340B policy change last year, beneficiaries are saving an estimated \$320 million on out-of-pocket payments for these drugs in 2018 alone.

The Food and Drug Administration's (FDA) <u>2017 generic drug approvals</u> contributed to prescription drug buyers saving \$8.8 billion. These savings were driven in part by the FDA hitting a record high of generic drug approvals last year.

The 2019 Part D <u>Explanation of Benefits</u> model will now allow plans to include information on drug price increases and more affordable formulary alternatives. The Administration is also including information for Part D beneficiaries on how to access the updated <u>CMS Drug Pricing Dashboards</u>.